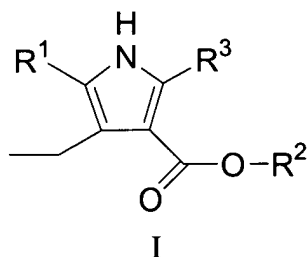


In the claims:

1. (Currently Amended) A compound of Formula I



wherein

R¹ is selected from

- 1) hydrogen,
- 2) halogen,
- 3) substituted or unsubstituted C₁-C₁₀ alkyl,
- 4) substituted or unsubstituted C₂-C₁₀ alkenyl,
- 5) substituted or unsubstituted C₂-C₁₀ alkynyl,
- 6) substituted or unsubstituted aryl,
- 7) substituted or unsubstituted C₃-C₁₀ cycloalkyl,
- 8) ~~substituted or unsubstituted heterocyclyl,~~
- 9) -(CR^{a2})_nOR⁴, and
- 10) -(CR^{a2})_tC(O)OR⁴;

said alkyl, alkenyl, alkynyl, aryl, and cycloalkyl, ~~and heterocyclyl~~ is optionally substituted with one or more of R⁷;

R² is selected from

- 1) hydrogen,
- 2) substituted or unsubstituted aralkyl,
- 3) substituted or unsubstituted C₁-C₁₀ alkyl,
- 4) ~~substituted or unsubstituted heterocyclyl,~~
- 5) substituted or unsubstituted aryl, and
- 6) substituted or unsubstituted C₃-C₁₀ cycloalkyl;

R³ is selected from

- 1) hydrogen,
- 2) halogen,
- 3) -C(O)R⁴,
- 4) substituted or unsubstituted C₁-C₁₀ alkyl,
- 5) substituted or unsubstituted aryl,
- 6) ~~substituted or unsubstituted heterocycetyl,~~
- 7) substituted or unsubstituted C₃-C₁₀ cycloalkyl,
- 8) substituted or unsubstituted C₂-C₁₀ alkenyl, and
- 9) substituted or unsubstituted C₂-C₁₀ alkynyl;

R⁴ is independently selected from

- 1) hydrogen,
- 2) substituted or unsubstituted C₁-C₁₀ alkyl,
- 3) substituted or unsubstituted aryl,
- 4) ~~substituted or unsubstituted heterocycetyl,~~
- 5) substituted or unsubstituted C₃-C₁₀ cycloalkyl,
- 6) substituted or unsubstituted C₂-C₁₀ alkenyl, and
- 7) substituted or unsubstituted C₂-C₁₀ alkynyl;

R⁶ is independently selected from

- 1) substituted or unsubstituted aryl,
- 2) ~~substituted or unsubstituted heterocycetyl,~~
- 3) substituted or unsubstituted cycloalkyl, and
- 4) halogen;

R⁷ is independently selected from

- 1) hydrogen,
- 2) halogen,
- 3) substituted or unsubstituted C₁-C₁₀ alkyl,
- 4) substituted or unsubstituted C₂-C₁₀ alkenyl,
- 5) substituted or unsubstituted C₂-C₁₀ alkynyl,

- 6) substituted or unsubstituted C₃-C₁₀ cycloalkyl,
- 7) substituted or unsubstituted aryl,
- ~~8) substituted or unsubstituted heterocyclyl,~~
- 9) -NO₂,
- 10) -NR⁴(CR^{a2})_nC(O)R⁴,
- 11) -(CR^{a2})_nNR⁴₂,
- 12) -(CR^{a2})_nNR⁴(CR^{a2})_nR⁶,
- 13) -CN,
- 14) -(CR^{a2})_nC(O)R⁴,
- 15) -(CR^{a2})_nC(O)(CR^{a2})_nOR⁴,
- 16) -(CR^{a2})_nOR⁴,
- 17) -(CR^{a2})_nR⁶,
- 18) -(CR^{a2})_nC(O)OR⁴, and
- 19) -(CR^{a2})_nSi(R⁴)₃;

R^a is independently selected from

- 1) hydrogen,
- 2) substituted or unsubstituted C₁-C₁₀ alkyl,
- 3) substituted or unsubstituted C₂-C₁₀ alkenyl,
- 4) substituted or unsubstitute C₂-C₁₀ alkynyl,
- 5) -OR⁴,
- 6) -C(O)OR⁴,
- 7) -NR⁴₂,
- 8) substituted or unsubstituted aryl,
- ~~9) substituted or unsubstituted heterocyclyl,~~ and
- 10) substituted or unsubstituted C₃-C₁₀ cycloalkyl;

n is independently 0 to 6;

t is 1 to 4;

or a pharmaceutically acceptable salt or stereoisomer thereof.

2. (Currently Amended) The compound according to Claim 1,
wherein

R¹ is selected from

- 1) hydrogen,
- 2) halogen,
- 3) substituted or unsubstituted C₁-C₆ alkyl,
- 4) substituted or unsubstituted C₂-C₁₀ alkynyl,
- 5) substituted or unsubstituted aryl, and
- 6) substituted or unsubstituted C₃-C₁₀ cycloalkyl, ~~and~~
- 7) ~~substituted or unsubstituted heterocyclyl;~~

said alkyl, alkynyl, aryl, and cycloalkyl, ~~and heterocyclyl~~ is optionally substituted with one or more of R⁷;

R² is selected from

- 1) substituted or unsubstituted aralkyl,
- 2) substituted or unsubstituted C₁-C₆ alkyl,
- 3) substituted or unsubstituted aryl, and
- 4) substituted or unsubstituted C₃-C₁₀ cycloalkyl;

R³ is selected from

- 1) halogen,
- 2) -C(O)R⁴, and
- 3) substituted or unsubstituted C₁-C₆ alkyl;

R⁴ is independently selected from

- 1) hydrogen,
- 2) substituted or unsubstituted C₁-C₆ alkyl,
- 1) substituted or unsubstituted aryl, and
- 2) ~~substituted or unsubstituted heterocyclyl, and~~
- 3) substituted or unsubstituted C₃-C₁₀ cycloalkyl;

or a pharmaceutically acceptable salt or stereoisomer thereof.

3. (Currently Amended) The compound according to Claim 2,

wherein

R¹ is selected from

- 1) substituted or unsubstituted C₁-C₆ alkyl,
- 2) substituted or unsubstituted C₂-C₁₀ alkynyl, and
- 3) ~~substituted or unsubstituted heterocyclyl and~~
- 4) substituted or unsubstituted aryl;

said alkyl, alkynyl, ~~heterocyclyl~~ and aryl is optionally substituted with one or more of R⁷;

R² is selected from

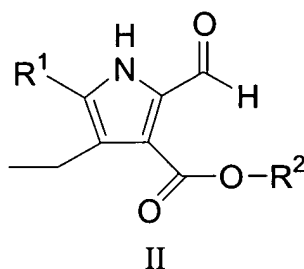
- 1) substituted or unsubstituted aralkyl, and
- 2) substituted or unsubstituted C₁-C₆ alkyl;

R³ is selected from

- 1) halogen, and
- 2) -C(O)R⁴;

or a pharmaceutically acceptable salt or stereoisomer thereof.

4. (Currently Amended) A compound of Formula II



wherein

R¹ is selected from

- 1) hydrogen,
- 2) halogen,
- 3) substituted or unsubstituted C₁-C₆ alkyl,
- 4) substituted or unsubstituted C₂-C₁₀ alkynyl,

- 5) substituted or unsubstituted aryl, and
- 6) substituted or unsubstituted C₃-C₁₀ cycloalkyl, and
- ~~7) substituted or unsubstituted heterocycyl;~~

said alkyl, alkynyl, aryl, and cycloalkyl ~~and heterocycyl~~ is optionally substituted with one or more of R⁷;

R² is selected from

- 1) substituted or unsubstituted aralkyl, and
- 2) substituted or unsubstituted C₁-C₆ alkyl;

R⁴ is independently selected from

- 1) hydrogen,
- 2) substituted or unsubstituted C₁-C₁₀ alkyl,
- 3) substituted or unsubstituted aryl,
- ~~4) substituted or unsubstituted heterocycyl;~~
- 5) substituted or unsubstituted C₃-C₁₀ cycloalkyl,
- 6) substituted or unsubstituted C₂-C₁₀ alkenyl, and
- 7) substituted or unsubstituted C₂-C₁₀ alkynyl;

R⁶ is independently selected from

- 1) substituted or ~~unsubstituted~~ unsubstituted aryl,
- ~~2) substituted or unsubstituted heterocycyl;~~
- 3) substituted or unsubstituted C₃-C₁₀ cycloalkyl, and
- 4) halogen;

R⁷ is independently selected from

- 1) hydrogen,
- 2) halogen,
- 3) substituted or unsubstituted C₁-C₁₀ alkyl,
- 4) substituted or unsubstituted C₂-C₁₀ alkenyl,
- 5) substituted or unsubstituted C₂-C₁₀ alkynyl,
- 6) substituted or unsubstituted C₃-C₁₀ cycloalkyl,
- 7) substituted or unsubstituted aryl,
- ~~8) substituted or unsubstituted heterocycyl;~~

- 9) $-\text{NO}_2$,
- 10) $-\text{NR}^4(\text{CRA}_2)_n\text{C}(\text{O})\text{R}^4$,
- 11) $-(\text{CRA}_2)_n\text{NR}^4_2$,
- 12) $-(\text{CRA}_2)_n\text{NR}^4(\text{CRA}_2)_n\text{R}^6$,
- 13) $-\text{CN}$,
- 14) $-(\text{CRA}_2)_n\text{C}(\text{O})\text{R}^4$,
- 15) $-(\text{CRA}_2)_n\text{C}(\text{O})(\text{CRA}_2)_n\text{OR}^4$,
- 16) $-(\text{CRA}_2)_n\text{OR}^4$,
- 17) $-(\text{CRA}_2)_n\text{R}^6$,
- 18) $-(\text{CRA}_2)_n\text{C}(\text{O})\text{OR}^4$, and
- 19) $-(\text{CRA}_2)_n\text{Si}(\text{R}^4)_3$;

R^a is independently selected from

- 1) hydrogen,
- 2) substituted or unsubstituted C_1 - C_{10} alkyl,
- 3) substituted or unsubstituted C_1 - C_{10} alkenyl,
- 4) substituted or unsubstituted C_1 - C_{10} alkynyl,
- 5) $-\text{OR}^4$,
- 6) $-\text{C}(\text{O})\text{OR}^4$,
- 7) $-\text{NR}^4_2$,
- 8) substituted or unsubstituted aryl, and
- 9) ~~substituted or unsubstituted heterocyclyl, and~~
- 10) substituted or unsubstituted C_3 - C_{10} cycloalkyl;

n is independently 0 to 6;

t is 1 to 4;

or a pharmaceutically acceptable salt or stereoisomer thereof.

5. (Currently Amended) A compound selected from:

benzyl 4-ethyl-2-formyl-1H-pyrrole-3-carboxylate;

benzyl 4-ethyl-2-formyl-5-iodo-1H-pyrrole-3-carboxylate;

methyl 4-ethyl-2-formyl-5-iodo-1H-pyrrole-3-carboxylate;

methyl 4-ethyl-2,5-diiodo-1H-pyrrole-3-carboxylate;

methyl 5-(4-fluorophenyl)-4-ethyl-2-formyl-1H-pyrrole-3-carboxylate;

methyl 4-ethyl-2-formyl-5-thien-2-yl-1H-pyrrole-3-carboxylate;

methyl 4-ethyl-2-formyl-5-[3-(trimethylsilyl)prop-1-ynyl]-1H-pyrrole-3-carboxylate;

4'-benzyl 1-tert-butyl 3'-ethyl-5'-formyl-1H,1'H-2,2'-bipyrrole-1,4'-dicarboxylate;

benzyl 5-(3,5-dimethylisoxazol-4-yl)-4-ethyl-2-formyl-1H-pyrrole-3-carboxylate;

benzyl 5-(1-benzofuran-2-yl)-4-ethyl-2-formyl-1H-pyrrole-3-carboxylate;

benzyl 4-ethyl-2-formyl-5-(3-nitrophenyl)-1H-pyrrole-3-carboxylate;

benzyl 4-ethyl-2-formyl-5-(5-methyl-2-furyl)-1H-pyrrole-3-carboxylate;

benzyl 5-[3-(acetylamino)phenyl]-4-ethyl-2-formyl-1H-pyrrole-3-carboxylate;

benzyl 4-ethyl-2-formyl-5-pyridin-4-yl-1H-pyrrole-3-carboxylate;

benzyl 4-ethyl-2-formyl-5-phenyl-1H-pyrrole-3-carboxylate;

benzyl 5-(3-cyanophenyl)-4-ethyl-2-formyl-1H-pyrrole-3-carboxylate;

benzyl 4-ethyl-2-formyl-5-(3-methoxyphenyl)-1H-pyrrole-3-carboxylate;

benzyl 4-ethyl-2-formyl-5-(5-formyl-2-furyl)-1H-pyrrole-3-carboxylate;

methyl 4-ethyl-2-formyl-5-(phenylethynyl)-1H-pyrrole-3-carboxylate;

methyl 5-{3-[benzyl(methyl)amino]prop-1-ynyl}-4-ethyl-2-formyl-1H-pyrrole-3-carboxylate;

benzyl 5-(2-cyanophenyl)-4-ethyl-2-formyl-1H-pyrrole-3-carboxylate;

benzyl 5-(4-cyanophenyl)-4-ethyl-2-formyl-1H-pyrrole-3-carboxylate;

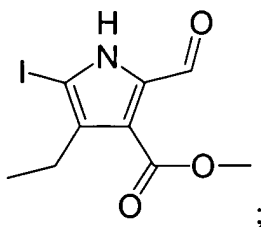
benzyl 4-ethyl-2-formyl-5-(4-nitrophenyl)-1H-pyrrole-3-carboxylate;

benzyl 4-ethyl-2-formyl-5-(2-methoxyphenyl)-1H-pyrrole-3-carboxylate;

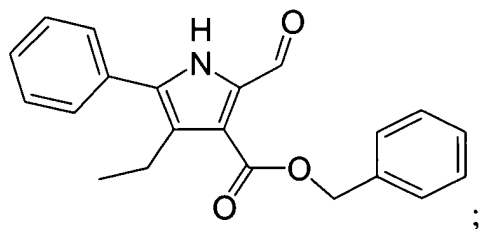
benzyl 4-ethyl-2-formyl-5-(4-methoxyphenyl)-1H-pyrrole-3-carboxylate;
benzyl 4-ethyl-2-formyl-5-(2-methylphenyl)-1H-pyrrole-3-carboxylate;
benzyl 4-ethyl-2-formyl-5-(3-methylphenyl)-1H-pyrrole-3-carboxylate;
benzyl 5-(2-chlorophenyl)-4-ethyl-2-formyl-1H-pyrrole-3-carboxylate;
benzyl 5-(3-chlorophenyl)-4-ethyl-2-formyl-1H-pyrrole-3-carboxylate;
methyl 4-ethyl-2-formyl-5-[1-(3-hydroxypropyl)vinyl]-1H-pyrrole-3-carboxylate;
methyl 4-ethyl-2-formyl-5-(5-hydroxypent-1-ynyl)-1H-pyrrole-3-carboxylate;
methyl 4-ethyl-2-formyl-5-[(1-hydroxycyclohexyl)ethynyl]-1H-pyrrole-3-carboxylate;
methyl 5-[3-(dimethylamino)prop-1-ynyl]-4-ethyl-2-formyl-1H-pyrrole-3-carboxylate;
methyl 5-(3,3-dimethylbut-1-ynyl)-4-ethyl-2-formyl-1H-pyrrole-3-carboxylate;
methyl 4-ethyl-2-formyl-5-(pyridin-2-ylethynyl)-1H-pyrrole-3-carboxylate;
methyl 4-ethyl-2-formyl-5-(6-methoxypyridin-2-yl)-1H-pyrrole-3-carboxylate;
methyl 4-ethyl-2-formyl-5-(3-methoxyprop-1-ynyl)-1H-pyrrole-3-carboxylate;
methyl 5-[(2-bromophenyl)ethynyl]-4-ethyl-2-formyl-1H-pyrrole-3-carboxylate;
methyl 5-[3-(1H-1,2,3-benzotriazol-1-yl)prop-1-ynyl]-4-ethyl-2-formyl-1H-pyrrole-3-carboxylate;
methyl 4-ethyl-5-(2-ethylbutyl)-2-formyl-1H-pyrrole-3-carboxylate;
methyl 4-ethyl-2-formyl-5-(4-methylpyridin-2-yl)-1H-pyrrole-3-carboxylate;
methyl 4-ethyl-2-formyl-5-(6-methylpyridin-2-yl)-1H-pyrrole-3-carboxylate;
methyl 5-(4-tert-butylphenyl)-4-ethyl-2-formyl-1H-pyrrole-3-carboxylate;
methyl 5-(2,4-difluorophenyl)-4-ethyl-2-formyl-1H-pyrrole-3-carboxylate;
methyl 4-ethyl-2-formyl-5-[3-(methoxycarbonyl)phenyl]-1H-pyrrole-3-carboxylate;
methyl 4-ethyl-2-formyl-5-[4-(methoxycarbonyl)phenyl]-1H-pyrrole-3-carboxylate;

methyl 4-ethyl-2-formyl-5-[(1-hydroxycyclopentyl)ethynyl]-1H-pyrrole-3-carboxylate;
methyl 4-ethyl-2-formyl-5-(3-hydroxy-3-methylbut-1-ynyl)-1H-pyrrole-3-carboxylate
methyl 4-ethyl-2-formyl-5-(1-hexylvinyl)-1H-pyrrole-3-carboxylate;
methyl 4-ethyl-2-formyl-5-(1,3-thiazol-2-yl)-1H-pyrrole-3-carboxylate;
methyl 5-[1-(3-chloropropyl)vinyl]-4-ethyl-2-formyl-1H-pyrrole-3-carboxylate;
methyl 5-(5-chloropent-1-ynyl)-4-ethyl-2-formyl-1H-pyrrole-3-carboxylate;
methyl 4-ethyl-2-formyl-5-(3-hydroxy-3-phenylbut-1-ynyl)-1H-pyrrole-3-carboxylate;
methyl 4-ethyl-2-formyl-5-(3-methylpyridin-2-yl)-1H-pyrrole-3-carboxylate;
methyl 4-ethyl-2-formyl-5-isopentyl-1H-pyrrole-3-carboxylate;
methyl 4-ethyl-2-formyl-5-(3-methylthien-2-yl)-1H-pyrrole-3-carboxylate;
methyl 4-ethyl-2-formyl-5-isobutyl-1H-pyrrole-3-carboxylate;
methyl 5-cyclohexyl-4-ethyl-2-formyl-1H-pyrrole-3-carboxylate;
methyl 5-butyl-4-ethyl-2-formyl-1H-pyrrole-3-carboxylate;
methyl 5-cyclopentyl-4-ethyl-2-formyl-1H-pyrrole-3-carboxylate;
methyl 5-(cyclohexylmethyl)-4-ethyl-2-formyl-1H-pyrrole-3-carboxylate;
methyl 5-sec-butyl-4-ethyl-2-formyl-1H-pyrrole-3-carboxylate;
methyl 4-ethyl-2-formyl-5-(3-methoxy-2-methyl-3-oxopropyl)-1H-pyrrole-3-carboxylate;
methyl 4-ethyl-2-formyl-5-phenyl-1H-pyrrole-3-carboxylate;
methyl 4-ethyl-2-formyl-5-pyridin-4-yl-1H-pyrrole-3-carboxylate;
methyl 4-ethyl-2-formyl-5-(4-nitrophenyl)-1H-pyrrole-3-carboxylate; and
methyl 4-ethyl-2-formyl-5-(2-methoxyphenyl)-1H-pyrrole-3-carboxylate;
or a pharmaceutically acceptable salt or stereoisomer thereof.

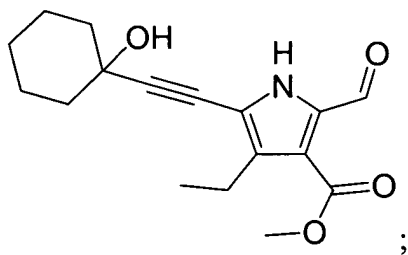
6. (Currently Amended) The compound according Claim 5 that is selected from
methyl 4-ethyl-2-formyl-5-iodo-1H-pyrrole-3-carboxylate



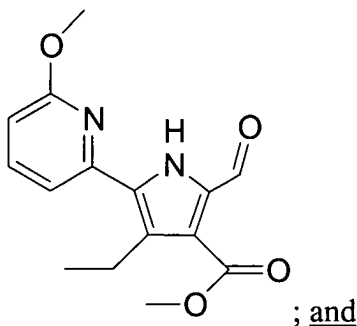
benzyl 4-ethyl-2-formyl-5-phenyl-1H-pyrrole-3-carboxylate



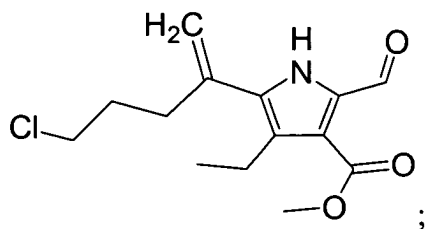
methyl 4-ethyl-2-formyl-5-[(1-hydroxycyclohexyl)ethynyl]-1H-pyrrole-3-carboxylate



methyl 4-ethyl-2-formyl-5-(6-methoxypyridin-2-yl)-1H-pyrrole-3-carboxylate



methyl 5-[1-(3-chloropropyl)vinyl]-4-ethyl-2-formyl-1H-pyrrole-3-carboxylate



or a pharmaceutically acceptable salt or stereoisomer thereof.

7. (Currently Amended) A trifluoroacetic acid salt of a compound of Claim 5 which is selected from

methyl 4-ethyl-2-formyl-5-(6-methoxypyridin-2-yl)-1H-pyrrole-3-carboxylate;

methyl 4-ethyl-2-formyl-5-(4-methylpyridin-2-yl)-1H-pyrrole-3-carboxylate;

methyl 4-ethyl-2-formyl-5-(6-methylpyridin-2-yl)-1H-pyrrole-3-carboxylate; and

benzyl 4-ethyl-2-formyl-5-pyridin-4-yl-1H-pyrrole-3-carboxylate.

8. (Original) A pharmaceutical composition which is comprised of a compound in accordance with Claim 1 and a pharmaceutically acceptable carrier.

9. (Original) A method of modulating the catalytic activity of protein kinases in a mammal in need thereof comprising contacting the protein kinase with a compound of Claim 1.

10. (Original) The method of Claim 9 wherein the protein kinase is an RTK.
11. (Original) The method of Claim 10, wherein the RTK is selected from IR, IGF-1R and IRR.
12. (Original) A method of treating a PK-related disorder in a mammal in need thereof comprising administering to said mammal a therapeutically effective amount of a compound of Claim 1.
13. (Currently Amended) A The method of Claim 12, wherein the PK-related disorder is an IGF-1R-related disorder selected from:
 - 1) cancer,
 - 2) diabetes,
 - 3) an autoimmune disorder,
 - 4) a hyperproliferation disorder,
 - 5) aging,
 - 6) acromegaly, and
 - 7) Crohn's disease.
14. (Cancelled)
15. (Cancelled)
16. (Original) A method of treating cancer in a mammal in need of such treatment comprising administering to said mammal a therapeutically effective amount of a compound of Claim 1.
17. (Original) A method of treating retinal vascularization comprising administering to a mammal in need of such treatment a therapeutically effective amount of a compound of Claim 1.
18. (Cancelled)

19. (Cancelled)

20. (Cancelled)

21. (Cancelled)

22. (Cancelled)

23. (Cancelled)

24. (Cancelled)

25. (Cancelled)

26. (Cancelled)

27. (Cancelled)

28. (Cancelled)

29. (Cancelled)

30. (Cancelled)